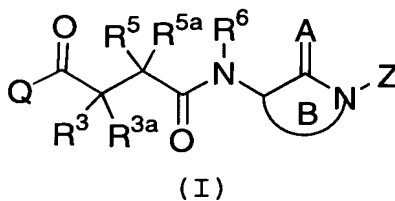


--1. (Twice Amended) A compound of Formula (I):



or a pharmaceutically acceptable salt thereof, wherein:

*Sub*  
*C1*  
A is O or S;

Q is  $-NR^1R^2$ ;

$R^1$  is selected from: H and  $C_1$ - $C_6$  alkyl;

$R^2$  is independently selected from H and  $C_1$ - $C_6$  alkyl;

$R^3$  is  $-(CR^7R^{7a})_n-R^4$ ,  
 $-(CR^7R^{7a})_n-S-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-O-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-N(R^{7b})-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-S(=O)-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-S(=O)_2-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-C(=O)-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-N(R^{7b})C(=O)-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-C(=O)N(R^{7b})-(CR^7R^{7a})_m-R^4$ ,  
 $-(CR^7R^{7a})_n-N(R^{7b})S(=O)_2-(CR^7R^{7a})_m-R^4$ , or  
 $-(CR^7R^{7a})_n-S(=O)_2N(R^{7b})-(CR^7R^{7a})_m-R^4$ ;

n is 0, 1, 2, or 3;

m is 0, 1, 2, or 3;

$R^{3a}$  is H, OH,  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  alkoxy,  $C_2$ - $C_4$  alkenyl  
or  $C_2$ - $C_4$  alkenyloxy;

B'  
Sub  
C1  
R<sup>4</sup> is H, OH, OR<sup>14a</sup>,

C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>6</sub> alkenyl substituted with 0-3 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>6</sub> alkynyl substituted with 0-3 R<sup>4a</sup>,  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;

R<sup>4a</sup>, at each occurrence, is independently selected from  
H, F, Cl, Br, I, CF<sub>3</sub>,  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,  
S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl,  
C<sub>1</sub>-C<sub>4</sub> haloalkoxy, and C<sub>1</sub>-C<sub>4</sub> haloalkyl-S-;

R<sup>5</sup> is H, OR<sup>14</sup>;

C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>1</sub>-C<sub>6</sub> alkoxy substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>6</sub> alkenyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>6</sub> alkynyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>5c</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>5c</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>;

*B*  
*Sub*  
*C1*  
~~R<sup>5a</sup> is H, OH, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>2</sub>-C<sub>4</sub> alkenyl, or C<sub>2</sub>-C<sub>4</sub> alkenyloxy;~~

~~R<sup>5b</sup>, at each occurrence, is independently selected from:  
H, C<sub>1</sub>-C<sub>6</sub> alkyl, CF<sub>3</sub>, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>,  
NR<sup>15</sup>R<sup>16</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>5c</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>5c</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>;~~

~~R<sup>5c</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,  
S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl,  
C<sub>1</sub>-C<sub>4</sub> haloalkoxy, and C<sub>1</sub>-C<sub>4</sub> haloalkyl-S-;~~

~~R<sup>6</sup> is H;  
C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>6a</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>6b</sup>; or  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>6b</sup>;~~

~~R<sup>6a</sup>, at each occurrence, is independently selected from H,  
C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>,  
aryl or CF<sub>3</sub>;~~

~~R<sup>6b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub>  
alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;~~

~~R<sup>7</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, CF<sub>3</sub>, phenyl and C<sub>1</sub>-C<sub>4</sub> alkyl;~~

*B*  
R<sup>7a</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, CF<sub>3</sub>, and C<sub>1</sub>-C<sub>4</sub> alkyl;

R<sup>7b</sup> is independently selected from H and C<sub>1</sub>-C<sub>4</sub> alkyl;

*Sub C*  
Ring B is a 7 membered lactam or thiolactam, wherein the lactam is 2-oxo-azepinyl or thiolactam is 2-thioxo-azepinyl;

wherein each additional lactam carbon or thiolactam carbon is substituted with 0-2 R<sup>11</sup>; provided two R<sup>11</sup> substituents are present on adjacent atoms and are combined to form a benzo fused radical; wherein said benzo fused radical is substituted with 0-4 R<sup>13</sup>;

and,

wherein the lactam or thiolactam contains a heteroatom selected from -N=, -NH-, and -N(R<sup>10</sup>)-;

R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, C(=O)NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>R<sup>17</sup>;

C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-3 R<sup>10a</sup>;

C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>10b</sup>;

C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or

5 to 10 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>10b</sup>;

R<sup>10a</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or aryl substituted with 0-4 R<sup>10b</sup>;

R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, and C<sub>1</sub>-C<sub>4</sub> haloalkyl-S-;

B1  
Sub  
C'  
R<sup>11</sup>, at each occurrence, is independently selected from  
H, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>18</sup>R<sup>19</sup>,  
C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, C(=O)NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>NR<sup>18</sup>R<sup>19</sup>, CF<sub>3</sub>;  
C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-3 R<sup>11a</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>11b</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>11b</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>11b</sup>;

R<sup>11a</sup>, at each occurrence, is independently selected from  
H, C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>,  
CF<sub>3</sub>;  
phenyl substituted with 0-3 R<sup>11b</sup>;  
C<sub>3</sub>-C<sub>6</sub> cycloalkyl substituted with 0-3 R<sup>11b</sup>; and  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>11b</sup>;

R<sup>11b</sup>, at each occurrence, is independently selected from H,  
OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,  
S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl,  
C<sub>1</sub>-C<sub>4</sub> haloalkoxy, and C<sub>1</sub>-C<sub>4</sub> haloalkyl-S-;

Z is H;

C<sub>1</sub>-C<sub>8</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>1</sub>-C<sub>8</sub> alkyl substituted with 0-3 R<sup>12a</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-4 R<sup>12b</sup>; or

B1  
Sub C1  
5 to 10 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>12b</sup>;

R<sup>12</sup>, at each occurrence, is independently selected from C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>; C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-4 R<sup>12b</sup>; or 5 to 10 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>12b</sup>;

R<sup>12a</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, -C(=O)NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, or C<sub>1</sub>-C<sub>4</sub> haloalkyl-S-;

R<sup>12b</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, and C<sub>1</sub>-C<sub>4</sub> haloalkyl-S-;

R<sup>13</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkoxyalkyl, or C<sub>3</sub>-C<sub>6</sub> cycloalkyl;

R<sup>14a</sup> is H, phenyl, benzyl, or C<sub>1</sub>-C<sub>4</sub> alkyl;

R<sup>15</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

*B'*  
R<sup>16</sup>, at each occurrence, is independently selected from  
H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, benzyl, phenethyl,  
(C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

*Sub C'*  
R<sup>17</sup> is H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkoxyalkyl,  
aryl substituted by 0-4 R<sup>17a</sup>, or  
-CH<sub>2</sub>-aryl substituted by 0-4 R<sup>17a</sup>;

R<sup>17a</sup> is H, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
propoxy, butoxy, -OH, F, Cl, Br, I, CF<sub>3</sub>, OCF<sub>3</sub>, SCH<sub>3</sub>,  
S(O)CH<sub>3</sub>, SO<sub>2</sub>CH<sub>3</sub>, -NH<sub>2</sub>, -N(CH<sub>3</sub>)<sub>2</sub>, or C<sub>1</sub>-C<sub>4</sub> haloalkyl;

R<sup>18</sup>, at each occurrence, is independently selected from  
H, C<sub>1</sub>-C<sub>6</sub> alkyl, phenyl, benzyl, phenethyl,  
(C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-; and

R<sup>19</sup>, at each occurrence, is independently selected from  
H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, phenyl, benzyl, phenethyl,  
(C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

provided, when R<sup>13</sup> is H,  
then Z is H;

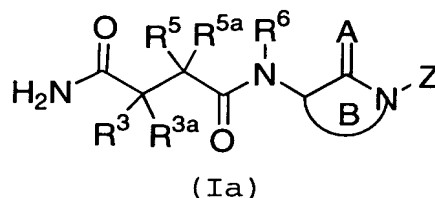
C<sub>4</sub>-C<sub>8</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>1</sub>-C<sub>8</sub> alkyl substituted with 0-3 R<sup>12a</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>; or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>; and

provided, when ring B is a 1,3,4,5-tetrahydro-1-(Z)-5-(R<sup>10</sup>)-  
6,6,7,7-tetra(R<sup>11</sup>)-2,4-dioxo-2H-1,5-diazepin-3-yl core, and  
R<sup>13</sup> is H; then

R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, C(=O)NR<sup>18</sup>R<sup>19</sup>,  
S(=O)<sub>2</sub>NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>R<sup>17</sup>; or  
C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-3 R<sup>10a</sup>;

Sub  
C1  
B1  
R<sup>10a</sup>, at each occurrence, is independently selected from  
H, C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>,  
and CF<sub>3</sub>.

Sub  
C2  
2. (Amended) A compound, according to Claim 1, of Formula  
(Ia):



or a pharmaceutically acceptable salt thereof,  
wherein:

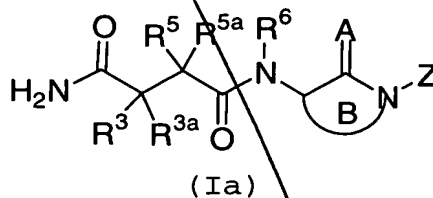
Z is H;

C<sub>1</sub>-C<sub>8</sub> alkyl substituted with 0-3 R<sup>12a</sup>;

C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>; or

C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>.

3. (Amended) A compound according to Claim 2 of Formula  
(Ia)



or a pharmaceutically acceptable salt thereof,  
wherein:

R<sup>3</sup> is -(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-R<sup>4</sup>,

-(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-S-(CR<sup>7</sup>R<sup>7a</sup>)<sub>m</sub>-R<sup>4</sup>,

-(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-O-(CR<sup>7</sup>R<sup>7a</sup>)<sub>m</sub>-R<sup>4</sup>, or

-(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-N(R<sup>7b</sup>)-(CR<sup>7</sup>R<sup>7a</sup>)<sub>m</sub>-R<sup>4</sup>;



n is 0, 1, or 2;

m is 0, 1, or 2;

R<sup>3a</sup> is H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, butoxy, allyl, or 3-buten-1-yl;

R<sup>4</sup> is H, OH, OR<sup>14a</sup>,

C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>4a</sup>,

C<sub>2</sub>-C<sub>6</sub> alkenyl substituted with 0-3 R<sup>4a</sup>,

C<sub>2</sub>-C<sub>6</sub> alkynyl substituted with 0-3 R<sup>4a</sup>,

C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,

C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or

5 to 10 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

sulphur, wherein said 5 to 10 membered heterocycle is

substituted with 0-3 R<sup>4b</sup>;

R<sup>4a</sup>, at each occurrence, is independently selected from

H, F, Cl, Br, I, CF<sub>3</sub>,

C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,

C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or

5 to 10 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

sulphur, wherein said 5 to 10 membered heterocycle is

substituted with 0-3 R<sup>4b</sup>;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,

Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,

S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub>

haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;

R<sup>5</sup> is H, OR<sup>14</sup>;

C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>5b</sup>,

C<sub>1</sub>-C<sub>6</sub> alkoxy substituted with 0-3 R<sup>5b</sup>,

C<sub>2</sub>-C<sub>6</sub> alkenyl substituted with 0-3 R<sup>5b</sup>,

C<sub>2</sub>-C<sub>6</sub> alkynyl substituted with 0-3 R<sup>5b</sup>;

B1  
C3-C10 carbocycle substituted with 0-3 R<sup>5c</sup>;  
C6-C10 aryl substituted with 0-3 R<sup>5c</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>;

Sub  
C2  
R<sup>5a</sup> is H or C<sub>1</sub>-C<sub>4</sub> alkyl;

R<sup>5b</sup>, at each occurrence, is independently selected from:  
H, C<sub>1</sub>-C<sub>6</sub> alkyl, CF<sub>3</sub>, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>,  
NR<sup>15</sup>R<sup>16</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>5c</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>5c</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>;

R<sup>5c</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,  
S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub>  
haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;

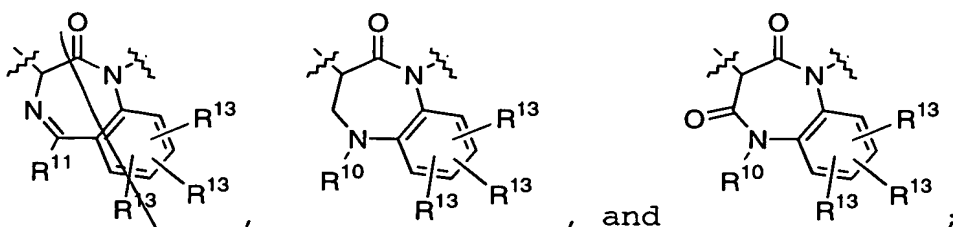
R<sup>6</sup> is H, methyl, or ethyl;

R<sup>7</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, CF<sub>3</sub>, phenyl and C<sub>1</sub>-C<sub>4</sub> alkyl;

R<sup>7a</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, CF<sub>3</sub>, and C<sub>1</sub>-C<sub>4</sub> alkyl;

R<sup>7b</sup> is independently selected from H, methyl, ethyl, propyl,  
and butyl;

Ring B is selected from



*Sub  
C2*

$R^{10}$  is H,  $C(=O)R^{17}$ ,  $C(=O)OR^{17}$ ,  $C(=O)NR^{18}R^{19}$ ,  
 $S(=O)_2NR^{18}R^{19}$ ,  $S(=O)_2R^{17}$ ;  
 $C_1$ - $C_6$  alkyl optionally substituted with 0-2  $R^{10a}$ ;  
 $C_6$ - $C_{10}$  aryl substituted with 0-4  $R^{10b}$ ;  
 $C_3$ - $C_{10}$  carbocycle substituted with 0-3  $R^{10b}$ ; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3  $R^{10b}$ ;

$R^{10a}$ , at each occurrence, is independently selected from H,  
 $C_1$ - $C_6$  alkyl,  $OR^{14}$ , Cl, F, Br, I, =O, CN,  $NO_2$ ,  $NR^{15}R^{16}$ ,  
 $CF_3$ , or phenyl substituted with 0-4  $R^{10b}$ ;

$R^{10b}$ , at each occurrence, is independently selected from H,  
OH,  $C_1$ - $C_6$  alkyl,  $C_1$ - $C_4$  alkoxy, Cl, F, Br, I, CN,  $NO_2$ ,  
 $NR^{15}R^{16}$ , or  $CF_3$ ;

$R^{11}$ , at each occurrence, is independently selected from  
H,  $C_1$ - $C_4$  alkoxy, Cl, F, Br, I, CN,  $NO_2$ ,  $NR^{18}R^{19}$ ,  $C(=O)R^{17}$ ,  
 $C(=O)OR^{17}$ ,  $C(=O)NR^{18}R^{19}$ ,  $S(=O)_2NR^{18}R^{19}$ ,  $CF_3$ ;  
 $C_1$ - $C_6$  alkyl optionally substituted with 0-3  $R^{11a}$ ;  
 $C_6$ - $C_{10}$  aryl substituted with 0-3  $R^{11b}$ ;  
 $C_3$ - $C_{10}$  carbocycle substituted with 0-3  $R^{11b}$ ; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3  $R^{11b}$ ;

Sub C2  
R<sup>11a</sup> at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-3 R<sup>11b</sup>;

R<sup>11b</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;

Z is H;

C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>12a</sup>;

C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>; or

C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>;

R<sup>12a</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, C<sub>1</sub>-C<sub>6</sub> alkyl, or C<sub>2</sub>-C<sub>6</sub> alkoxyalkyl;

R<sup>14a</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

R<sup>15</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

R<sup>16</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

R<sup>17</sup> is H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkoxyalkyl, aryl substituted by 0-4 R<sup>17a</sup>, or

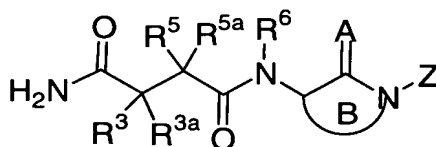
~~-CH<sub>2</sub>-aryl substituted by 0-4 R<sup>17a</sup>;~~

~~R<sup>17a</sup> is H, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, butoxy, -OH, F, Cl, Br, I, CF<sub>3</sub>, OCF<sub>3</sub>, SCH<sub>3</sub>, S(O)CH<sub>3</sub>, SO<sub>2</sub>CH<sub>3</sub>, -NH<sub>2</sub>, -N(CH<sub>3</sub>)<sub>2</sub>, or C<sub>1</sub>-C<sub>4</sub> haloalkyl;~~

~~R<sup>18</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, phenyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-; and~~

~~R<sup>19</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, phenyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-.~~

~~4. (Twice Amended) A compound according to Claim 3 of Formula (Ia)~~



~~(Ia)~~

~~or a pharmaceutically acceptable salt thereof,  
wherein:~~

~~R<sup>3</sup> is -(CHR<sup>7</sup>)<sub>n</sub>-R<sup>4</sup>,~~

~~n is 0 or 1;~~

~~R<sup>3a</sup> is H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, butoxy, allyl, or 3-buten-1-yl;~~

~~R<sup>4</sup> is H, OH, OR<sup>14a</sup>,~~

~~C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-2 R<sup>4a</sup>,~~

~~C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-2 R<sup>4a</sup>,~~

~~C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>,~~

~~B1~~  
~~Sub C3~~  
~~C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;~~

~~R<sup>4a</sup>, at each occurrence, is independently selected from  
H, F, Cl, Br, I, CF<sub>3</sub>,  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
phenyl substituted with 0-3 R<sup>4b</sup>, or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;~~

~~R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,  
S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub>  
haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;~~

~~R<sup>5</sup> is H, OR<sup>14</sup>;  
C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>5b</sup>;~~

~~R<sup>5a</sup> is H, methyl, ethyl, propyl, or butyl;~~

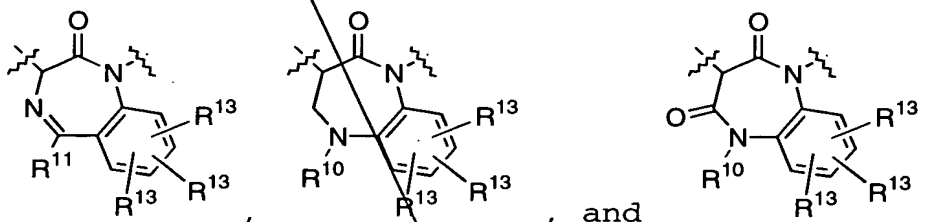
~~R<sup>5b</sup>, at each occurrence, is independently selected from:  
H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, Cl, F, Br, I,  
=O;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>5c</sup>;  
phenyl substituted with 0-3 R<sup>5c</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>;~~

B1  
Sub C3  
R<sup>5c</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>6</sup> is H;

R<sup>7</sup>, at each occurrence, is independently selected from H, F, CF<sub>3</sub>, methyl, and ethyl;

Ring B is selected from



R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>;  
C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>10a</sup>;  
phenyl substituted with 0-4 R<sup>10b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>10b</sup>;

R<sup>10a</sup> is selected from H, C<sub>1</sub>-C<sub>4</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-4 R<sup>10b</sup>;

R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, or CF<sub>3</sub>;

R<sup>11</sup> is selected from

B1  
Sub  
C3  
~~H, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, NR<sup>18</sup>R<sup>19</sup>, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, CF<sub>3</sub>;  
C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-3 R<sup>11a</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>11b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>11b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>11b</sup>;~~

R<sup>11a</sup>, at each occurrence, is independently selected from H,  
C<sub>1</sub>-C<sub>4</sub> alkyl, OR<sup>14</sup>, F, =O, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl  
substituted with 0-3 R<sup>11b</sup>;

R<sup>11b</sup>, at each occurrence, is independently selected from H,  
OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub>  
haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

Z is H;

C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-3 R<sup>12a</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>; or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>;

R<sup>12a</sup>, at each occurrence, is independently selected from  
H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>,  
S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and  
C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from  
H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>,  
NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, C<sub>1</sub>-C<sub>4</sub> alkyl, or C<sub>2</sub>-C<sub>4</sub> alkoxyalkyl;

R<sup>15</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-  
C<sub>4</sub> alkyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>4</sub> alkyl)-C(=O)-, and  
(C<sub>1</sub>-C<sub>4</sub> alkyl)-S(=O)<sub>2</sub>-;



~~R<sup>16</sup>, at each occurrence, is independently selected from  
H, OH, C<sub>1</sub>-C<sub>4</sub> alkyl, benzyl, phenethyl,  
(C<sub>1</sub>-C<sub>4</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>4</sub> alkyl)-S(=O)<sub>2</sub>-;~~

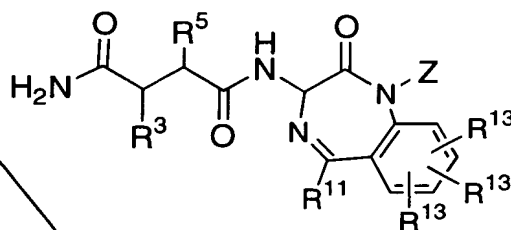
~~R<sup>17</sup> is H, methyl, ethyl, propyl, butyl, methoxymethyl,  
ethoxymethyl, methoxyethyl, ethoxyethyl,  
phenyl substituted by 0-3 R<sup>17a</sup>, or  
-CH<sub>2</sub>-phenyl substituted by 0-3 R<sup>17a</sup>;~~

~~R<sup>17a</sup> is H, methyl, methoxy, -OH, F, Cl, CF<sub>3</sub>, or OCF<sub>3</sub>;~~

~~R<sup>18</sup>, at each occurrence, is independently selected from  
H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and  
phenethyl; and~~

~~R<sup>19</sup>, at each occurrence, is independently selected from  
H, methyl, and ethyl.~~

~~6. (Twice Amended) A compound according to Claim 4 of Formula  
(Ic):~~



~~(Ic)~~

~~or a pharmaceutically acceptable salt thereof  
wherein~~

~~R<sup>3</sup> is R<sup>4</sup>,~~

~~R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>4a</sup>, or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>;~~

*B2*  
*Sub*  
*C4*  
R<sup>4a</sup> is selected from

H, F, CF<sub>3</sub>,

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,

phenyl substituted with 0-3 R<sup>4b</sup>, or

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

sulphur, wherein said 5 to 6 membered heterocycle is

substituted with 0-3 R<sup>4b</sup>; wherein said 5 to 6 membered

heterocycle is selected from pyridinyl, pyrimidinyl,

triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,

piperazinyl, piperidinyl, pyrazolyl, imidazolyl,

oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>5</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>5b</sup>;

R<sup>5b</sup> is selected from:

H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, =O;

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-2 R<sup>5c</sup>;

phenyl substituted with 0-3 R<sup>5c</sup>; or

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

sulphur, wherein said 5 to 6 membered heterocycle is

substituted with 0-3 R<sup>5c</sup>; wherein said 5 to 6 membered

heterocycle is selected from pyridinyl, pyrimidinyl,

triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,

piperazinyl, piperidinyl, pyrazolyl, imidazolyl,

oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>5c</sup>, at each occurrence, is independently selected from H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,

methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy,  
C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>11</sup> is selected from

H, NR<sup>18</sup>R<sup>19</sup>, CF<sub>3</sub>;

C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>11a</sup>;

phenyl substituted with 0-3 R<sup>11b</sup>;

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>11b</sup>; and

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>11b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>11a</sup> is selected from H, C<sub>1</sub>-C<sub>4</sub> alkyl, OR<sup>14</sup>, F, =O, NR<sup>15</sup>R<sup>16</sup>,  
CF<sub>3</sub>, or phenyl substituted with 0-3 R<sup>11b</sup>;

R<sup>11b</sup>, at each occurrence, is independently selected from H,  
OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, methyl, ethyl, propyl, butyl,  
methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub>  
haloalkoxy;

Z is H;

C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-3 R<sup>12a</sup>;

C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>; or

C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>;

R<sup>12a</sup>, at each occurrence, is independently selected from  
H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>,  
S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from

H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, Cl, F, Br, CN, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

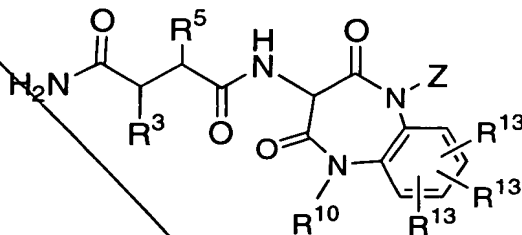
R<sup>15</sup>, at each occurrence, is independently selected from H, methyl, ethyl, propyl, and butyl;

R<sup>16</sup>, at each occurrence, is independently selected from H, OH, methyl, ethyl, propyl, butyl, benzyl, phenethyl, methyl-C(=O)-, ethyl-C(=O)-, methyl-S(=O)<sub>2</sub>-, and ethyl-S(=O)<sub>2</sub>-;

R<sup>18</sup>, at each occurrence, is independently selected from H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and phenethyl; and

R<sup>19</sup>, at each occurrence, is independently selected from H, methyl, and ethyl.

8. (Twice Amended) A compound according to Claim 4 of Formula (Ie):



(Ie)

or a pharmaceutically acceptable salt thereof wherein:

R<sup>3</sup> is R<sup>4</sup>,

R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>4a</sup>, or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>;

~~R<sup>4a</sup> is selected from~~

~~H, F, CF<sub>3</sub>,~~

~~C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,~~

~~phenyl substituted with 0-3 R<sup>4b</sup>, or~~

~~5 to 6 membered heterocycle containing 1 to 4~~

~~heteroatoms selected from nitrogen, oxygen, and~~

~~sulphur, wherein said 5 to 6 membered heterocycle is~~

~~substituted with 0-3 R<sup>4b</sup>; wherein said 5 to 6 membered~~

~~heterocycle is selected from pyridinyl, pyrimidinyl,~~

~~triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,~~

~~piperazinyl, piperidinyl, pyrazolyl, imidazolyl,~~

~~oxazolyl, isoxazolyl, and tetrazolyl;~~

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>5</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>5b</sup>;

R<sup>5b</sup> is selected from:

H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, =O;

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-2 R<sup>5c</sup>;

phenyl substituted with 0-3 R<sup>5c</sup>; or

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

sulphur, wherein said 5 to 6 membered heterocycle is

substituted with 0-3 R<sup>5c</sup>; wherein said 5 to 6 membered

heterocycle is selected from pyridinyl, pyrimidinyl,

triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,

piperazinyl, piperidinyl, pyrazolyl, imidazolyl,

oxazolyl, isoxazolyl, and tetrazolyl;

B3  
R<sup>5c</sup>, at each occurrence, is independently selected from H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

Sub  
CS  
R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>;  
C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>10a</sup>;  
phenyl substituted with 0-4 R<sup>10b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>10b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>10a</sup> is selected from H, methyl, ethyl, propyl, butyl, OR<sup>14</sup>, Cl, F, =O, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-4 R<sup>10b</sup>;

R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, Cl, F, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

Z is H;

C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-3 R<sup>12a</sup>;

C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>; or

C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>;

R<sup>12a</sup>, at each occurrence, is independently selected from H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from

H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
Cl, F, Br, CN, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

*B2*  
R<sup>14</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

R<sup>15</sup>, at each occurrence, is independently selected from H,  
methyl, ethyl, propyl, and butyl;

*Sub*  
*CS*  
R<sup>16</sup>, at each occurrence, is independently selected from  
H, OH, methyl, ethyl, propyl, butyl, benzyl, phenethyl,  
methyl-C(=O)-, ethyl-C(=O)-,  
methyl-S(=O)<sub>2</sub>-, and ethyl-S(=O)<sub>2</sub>-;

R<sup>17</sup> is H, methyl, ethyl, propyl, butyl, methoxymethyl,  
ethoxymethyl, methoxyethyl, ethoxyethyl,  
phenyl substituted by 0-3 R<sup>17a</sup>, or  
-CH<sub>2</sub>-phenyl substituted by 0-3 R<sup>17a</sup>;

R<sup>17a</sup> is H, methyl, methoxy, -OH, F, Cl, CF<sub>3</sub>, or OCF<sub>3</sub>;

R<sup>18</sup>, at each occurrence, is independently selected from  
H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and  
phenethyl; and

R<sup>19</sup>, at each occurrence, is independently selected from  
H, methyl, and ethyl.

*B2*  
*Sub*  
*C6*  
10. (Twice Amended) A compound, according to one of Claims 6,  
8, or 25 wherein:

R<sup>3</sup> is -CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>,  
-CH(CH<sub>3</sub>)<sub>2</sub>, -CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH(CH<sub>3</sub>)<sub>2</sub>,  
-CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>,  
-CH=CH<sub>2</sub>, -CH<sub>2</sub>CH=CH<sub>2</sub>, -CH<sub>2</sub>C(CH<sub>3</sub>)=CH<sub>2</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH=CH<sub>2</sub>,  
cis-CH<sub>2</sub>CH=CH(CH<sub>3</sub>),

*B \$*  
*Sub*  
*C6*

~~trans-CH<sub>2</sub>CH=CH(CH<sub>3</sub>),  
-C≡CH, -CH<sub>2</sub>C≡CH, -CH<sub>2</sub>C≡C(CH<sub>3</sub>),  
cyclopropyl-CH<sub>2</sub>-, cyclobutyl-CH<sub>2</sub>-, cyclopentyl-CH<sub>2</sub>-,  
cyclohexyl-CH<sub>2</sub>-, cyclopropyl-CH<sub>2</sub>CH<sub>2</sub>-,  
cyclobutyl-CH<sub>2</sub>CH<sub>2</sub>-, cyclopentyl-CH<sub>2</sub>CH<sub>2</sub>-,  
cyclohexyl-CH<sub>2</sub>CH<sub>2</sub>-, phenyl-CH<sub>2</sub>-,  
(2-F-phenyl)CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>-, (4-F-phenyl)CH<sub>2</sub>-,  
(2-Cl-phenyl)CH<sub>2</sub>-, (3-Cl-phenyl)CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>-,  
(2,3-diF-phenyl)CH<sub>2</sub>-, (2,4-diF-phenyl)CH<sub>2</sub>-,  
(2,5-diF-phenyl)CH<sub>2</sub>-, (2,6-diF-phenyl)CH<sub>2</sub>-,  
(3,4-diF-phenyl)CH<sub>2</sub>-, (3,5-diF-phenyl)CH<sub>2</sub>-,  
(2,3-diCl-phenyl)CH<sub>2</sub>-, (2,4-diCl-phenyl)CH<sub>2</sub>-,  
(2,5-diCl-phenyl)CH<sub>2</sub>-, (2,6-diCl-phenyl)CH<sub>2</sub>-,  
(3,4-diCl-phenyl)CH<sub>2</sub>-, (3,5-diCl-phenyl)CH<sub>2</sub>-,  
(3-F-4-Cl-phenyl)CH<sub>2</sub>-, (3-F-5-Cl-phenyl)CH<sub>2</sub>-,  
(3-Cl-4-F-phenyl)CH<sub>2</sub>-, phenyl-CH<sub>2</sub>CH<sub>2</sub>-,  
(2-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,3-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,4-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,5-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,6-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3,4-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3,5-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,3-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,4-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,5-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,6-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3,4-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3,5-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-F-4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, or (3-F-5-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,~~

R<sup>5</sup> is -CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH(CH<sub>3</sub>)<sub>2</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>,  
-CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH(CH<sub>3</sub>)<sub>2</sub>, -CH<sub>2</sub>C(CH<sub>3</sub>)<sub>3</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>3</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH(CH<sub>3</sub>)<sub>2</sub>, -CH(CH<sub>2</sub>CH<sub>3</sub>)<sub>2</sub>, -CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>, -CH=CH<sub>2</sub>, -CH<sub>2</sub>CH=CH<sub>2</sub>,  
-CH=CHCH<sub>3</sub>, cis-CH<sub>2</sub>CH=CH(CH<sub>3</sub>), trans-CH<sub>2</sub>CH=CH(CH<sub>3</sub>),  
trans-CH<sub>2</sub>CH=CH(C<sub>6</sub>H<sub>5</sub>), -CH<sub>2</sub>CH=C(CH<sub>3</sub>)<sub>2</sub>, cis-CH<sub>2</sub>CH=CHCH<sub>2</sub>CH<sub>3</sub>,  
trans-CH<sub>2</sub>CH=CHCH<sub>2</sub>CH<sub>3</sub>, cis-CH<sub>2</sub>CH<sub>2</sub>CH=CH(CH<sub>3</sub>),  
trans-CH<sub>2</sub>CH<sub>2</sub>CH=CH(CH<sub>3</sub>), trans-CH<sub>2</sub>CH=CHCH<sub>2</sub>(C<sub>6</sub>H<sub>5</sub>),  
-C≡CH, -CH<sub>2</sub>C≡CH, -CH<sub>2</sub>C≡C(CH<sub>3</sub>), -CH<sub>2</sub>C≡C(C<sub>6</sub>H<sub>5</sub>),



B\$  
Sub  
C6

~~-CH<sub>2</sub>CH<sub>2</sub>C≡CH, -CH<sub>2</sub>CH<sub>2</sub>C≡C(CH<sub>3</sub>), -CH<sub>2</sub>CH<sub>2</sub>C≡C(C<sub>6</sub>H<sub>5</sub>),  
cyclopropyl-CH<sub>2</sub>-, cyclobutyl-CH<sub>2</sub>-, cyclopentyl-CH<sub>2</sub>-,  
cyclohexyl-CH<sub>2</sub>-, (2-CH<sub>3</sub>-cyclopropyl)CH<sub>2</sub>-,  
(3-CH<sub>3</sub>-cyclobutyl)CH<sub>2</sub>-,  
cyclopropyl-CH<sub>2</sub>CH<sub>2</sub>-, cyclobutyl-CH<sub>2</sub>CH<sub>2</sub>-,  
cyclopentyl-CH<sub>2</sub>CH<sub>2</sub>-, cyclohexyl-CH<sub>2</sub>CH<sub>2</sub>-,  
(2-CH<sub>3</sub>-cyclopropyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-CH<sub>3</sub>-cyclobutyl)CH<sub>2</sub>CH<sub>2</sub>-,  
phenyl-CH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>-,  
(4-F-phenyl)CH<sub>2</sub>-, furanyl-CH<sub>2</sub>-, thienyl-CH<sub>2</sub>-,  
pyridyl-CH<sub>2</sub>-, 1-imidazolyl-CH<sub>2</sub>-, oxazolyl-CH<sub>2</sub>-,  
isoxazolyl-CH<sub>2</sub>-,  
phenyl-CH<sub>2</sub>CH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, furanyl-CH<sub>2</sub>CH<sub>2</sub>-, thienyl-CH<sub>2</sub>CH<sub>2</sub>-,  
pyridyl-CH<sub>2</sub>CH<sub>2</sub>-, 1-imidazolyl-CH<sub>2</sub>CH<sub>2</sub>-, oxazolyl-CH<sub>2</sub>CH<sub>2</sub>-,  
isoxazolyl-CH<sub>2</sub>CH<sub>2</sub>-;~~

Z is methyl, ethyl, i-propyl, n-propyl, n-butyl, i-butyl, s-butyl, t-butyl, or allyl;

R<sup>10</sup> is H, methyl, ethyl, phenyl, benzyl, phenethyl,  
4-F-phenyl, (4-F-phenyl)CH<sub>2</sub>-, (4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-Cl-phenyl, (4-Cl-phenyl)CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CH<sub>3</sub>-phenyl, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>-, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CF<sub>3</sub>-phenyl, (4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>-, or  
(4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-;

R<sup>11</sup>, at each occurrence, is independently selected from  
H, =O, methyl, ethyl, phenyl, benzyl, phenethyl,  
4-F-phenyl, (4-F-phenyl)CH<sub>2</sub>-, (4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
3-F-phenyl, (3-F-phenyl)CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
2-F-phenyl, (2-F-phenyl)CH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-Cl-phenyl, (4-Cl-phenyl)CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
3-Cl-phenyl, (3-Cl-phenyl)CH<sub>2</sub>-, (3-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CH<sub>3</sub>-phenyl, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>-, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
3-CH<sub>3</sub>-phenyl, (3-CH<sub>3</sub>-phenyl)CH<sub>2</sub>-, (3-CH<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CF<sub>3</sub>-phenyl, (4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>-, (4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
pyrid-2-yl, pyrid-3-yl, or pyrid-4-yl, and

Sub  
C6

R<sup>13</sup>, at each occurrence, is independently selected from  
H, F, Cl, OH, -CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>3</sub>, -OCH<sub>3</sub>, or -CF<sub>3</sub>.

B# 11. (Amended) A compound according to Claim 2 selected from:

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

Sub  
C7

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-propyl-butanediamide;

(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-propyl-butanediamide;

(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-propyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-methyl-3-allyl-butanediamide;

(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-methyl-3-allyl-butanediamide;

(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-methyl-3-propyl-butanediamide;

(2R) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-methyl-butanediamide;

*Sub C7*  
~~(2R,3S) N1-[1,3-dihydro-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(2-fluorophenyl)-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(2-fluorophenyl)-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-(2-fluorophenyl)-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2S,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-propyl-butanediamide;~~

(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(2-fluorophenyl)-7-chloro-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-propyl-butanediamide;

~~BA~~  
(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(4-fluorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

~~Sub C7~~  
(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(4-fluorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-(4-fluorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(pyrid-2-yl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(N-morpholino)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(dimethylamino)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(N-methyl-N-phenylamino)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(N-piperidinyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(N-homopiperidinyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

34  
(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(3-methoxyphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

Sub  
C7  
(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(pyrid-4-yl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-7-methoxy-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(pyrid-3-yl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-(cyclopropylmethyl)-butanediamide;

(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(3-fluorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(3-fluorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-(3-fluorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;

(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-(3-buten-1-yl)-butanediamide;

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-(cyclopentylethyl)-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(4-trifluoromethylphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-(3-buten-1-yl)-butanediamide;~~

~~(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-(4-trifluoromethylphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-(3-buten-1-yl)-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(4-trifluoromethylphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(4-trifluoromethylphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3R)-1,3-dihydro-1-methyl-2-oxo-5-(4-trifluoromethylphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(4-trifluoromethylphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-n-butyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(4-trifluoromethylphenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-propyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(4-chlorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-(3-buten-1-yl)-butanediamide;~~

BA  
Sub  
C7  
~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-(4-chlorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-n-butyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-N4-[benzyl]-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-methyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-n-butyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(2-methylpropyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-methyl-2-oxo-5-(4-chlorophenyl)-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-ethyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

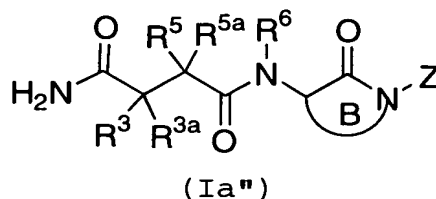
~~(2R,3S) N1-[1,3-dihydro-1-propyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[1,3-dihydro-1-(isopropyl)-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide;~~

~~(2R,3S) N1-[(3S)-1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl]-2-(2-methylpropyl)-3,3-diallyl-butanediamide; and~~

~~(2R,3S) N1-[1,3,4,5-tetrahydro-1,5-dimethyl-2,4-dioxo-2H-1,5-benzodiazepin-3-yl]-2-(2-methylpropyl)-3-allyl-butanediamide.~~

BA  
12. (Amended) A compound, according to Claim 1, of Formula (Ia''):



or a pharmaceutically acceptable salt thereof,  
wherein:

Z is C<sub>1</sub>-C<sub>8</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-4 R<sup>12b</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>12b</sup>;

provided, when R<sup>13</sup> is H,  
then Z is C<sub>4</sub>-C<sub>8</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>; or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>; and

provided, when ring B is a 1,3,4,5-tetrahydro-1-(Z)-5-(R<sup>10</sup>)-  
6,6,7,7-tetra(R<sup>11</sup>)-2,4-dioxo-2H-1,5-diazepin-3-yl core, and  
R<sup>13</sup> is H; then

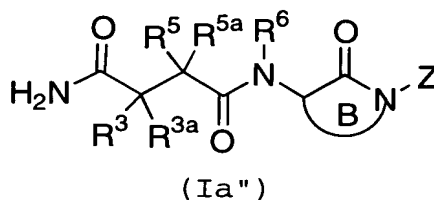
R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, C(=O)NR<sup>18</sup>R<sup>19</sup>,  
S(=O)<sub>2</sub>NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>R<sup>17</sup>; or  
C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-3 R<sup>10a</sup>; and

R<sup>10a</sup>, at each occurrence, is independently selected from



H, C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>.

**13.** (Amended) A compound according to Claim 12 of Formula (Ia")



or a pharmaceutically acceptable salt thereof,  
wherein:

R<sup>3</sup> is -(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-R<sup>4</sup>,  
-(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-S-(CR<sup>7</sup>R<sup>7a</sup>)<sub>m</sub>-R<sup>4</sup>,  
-(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-O-(CR<sup>7</sup>R<sup>7a</sup>)<sub>m</sub>-R<sup>4</sup>, or  
-(CR<sup>7</sup>R<sup>7a</sup>)<sub>n</sub>-N(R<sup>7b</sup>)-(CR<sup>7</sup>R<sup>7a</sup>)<sub>m</sub>-R<sup>4</sup>;

n is 0, 1, or 2;

m is 0, 1, or 2;

R<sup>3a</sup> is H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, butoxy, allyl, or 3-buten-1-yl;

R<sup>4</sup> is H, OH, OR<sup>14a</sup>,  
C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>6</sub> alkenyl substituted with 0-3 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>6</sub> alkynyl substituted with 0-3 R<sup>4a</sup>,  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;

Sub C7  
R<sup>4a</sup>, at each occurrence, is independently selected from  
H, F, Cl, Br, I, CF<sub>3</sub>,  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,  
S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub>  
haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;

R<sup>5</sup> is H, OR<sup>14</sup>;  
C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>1</sub>-C<sub>6</sub> alkoxy substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>6</sub> alkenyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>6</sub> alkynyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>5c</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>5c</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>;

R<sup>5a</sup> is H or C<sub>1</sub>-C<sub>4</sub> alkyl;

R<sup>5b</sup>, at each occurrence, is independently selected from:  
H, C<sub>1</sub>-C<sub>6</sub> alkyl, CF<sub>3</sub>, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>,  
NR<sup>15</sup>R<sup>16</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>5c</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>5c</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and

~~sub~~ sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>5c</sup>;

~~R<sup>5c</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;~~

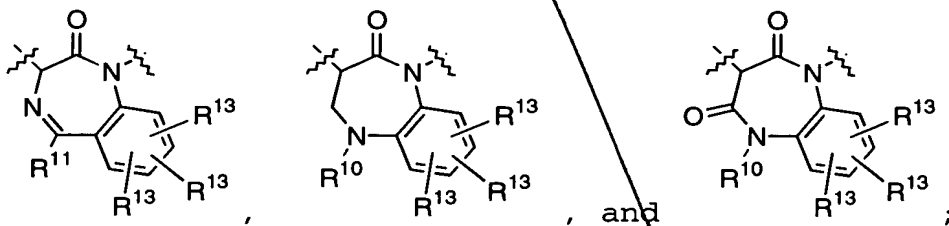
~~R<sup>6</sup> is H, methyl, or ethyl;~~

~~R<sup>7</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, CF<sub>3</sub>, phenyl, and C<sub>1</sub>-C<sub>4</sub> alkyl;~~

~~R<sup>7a</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, CF<sub>3</sub>, and C<sub>1</sub>-C<sub>4</sub> alkyl;~~

~~R<sup>7b</sup> is independently selected from H, methyl, ethyl, propyl, and butyl;~~

~~Ring B is selected from~~



~~R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, C(=O)NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>R<sup>17</sup>;~~

~~C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-2 R<sup>10a</sup>;~~

~~C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>10b</sup>;~~

~~C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or~~

~~5 to 10 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>10b</sup>;~~

B#  
R<sup>10a</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-4 R<sup>10b</sup>;

R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, or CF<sub>3</sub>;

Sub  
C7  
R<sup>11</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>18</sup>R<sup>19</sup>, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, C(=O)NR<sup>18</sup>R<sup>19</sup>, S(=O)<sub>2</sub>NR<sup>18</sup>R<sup>19</sup>, CF<sub>3</sub>; C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-3 R<sup>11a</sup>; C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>11b</sup>; C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-3 R<sup>11b</sup>; or 5 to 10 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>11b</sup>;

R<sup>11a</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-3 R<sup>11b</sup>;

R<sup>11b</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;

Z is C<sub>1</sub>-C<sub>6</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-4 R<sup>12b</sup>; or  
5 to 10 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and

BA  
Suh  
C7  
sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>12b</sup>;

R<sup>12</sup>, at each occurrence, is independently selected from C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>; C<sub>3</sub>-C<sub>10</sub> carbocycle substituted with 0-4 R<sup>12b</sup>; or 5 to 10 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 10 membered heterocycle is substituted with 0-3 R<sup>12b</sup>;

R<sup>12b</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkyl, and C<sub>1</sub>-C<sub>4</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, C<sub>1</sub>-C<sub>6</sub> alkyl, or C<sub>2</sub>-C<sub>6</sub> alkoxyalkyl;

R<sup>14a</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

R<sup>15</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

R<sup>16</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

R<sup>17</sup> is H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkoxyalkyl, aryl substituted by 0-4 R<sup>17a</sup>, or -CH<sub>2</sub>-aryl substituted by 0-4 R<sup>17a</sup>;

*BA*  
*Sub*  
*c7*  
R<sup>17a</sup> is H, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, butoxy, -OH, F, Cl, Br, I, CF<sub>3</sub>, OCF<sub>3</sub>, SCH<sub>3</sub>, S(O)CH<sub>3</sub>, SO<sub>2</sub>CH<sub>3</sub>, -NH<sub>2</sub>, -N(CH<sub>3</sub>)<sub>2</sub>, or C<sub>1</sub>-C<sub>4</sub> haloalkyl;

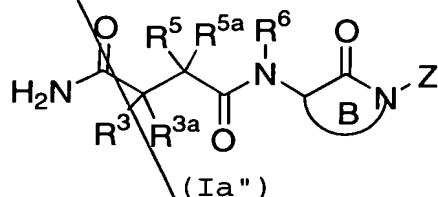
R<sup>18</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, phenyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-; and

R<sup>19</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, phenyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>6</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>6</sub> alkyl)-S(=O)<sub>2</sub>-;

provided, when R<sup>13</sup> is H,

then Z is C<sub>4</sub>-C<sub>6</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>; or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>.

**14.** (Amended) A compound according to Claim 13 of Formula (Ia")



or a pharmaceutically acceptable salt thereof,  
wherein:

R<sup>3</sup> is -(CHR<sup>7</sup>)<sub>n</sub>-R<sup>4</sup>,

n is 0 or 1;

R<sup>3a</sup> is H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, butoxy, allyl, or 3-buten-1-yl;

R<sup>4</sup> is H, OH, OR<sup>14a</sup>,

C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-2 R<sup>4a</sup>,

BA  
Sub  
C7  
~~C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-2 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>,  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>4b</sup>, or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;~~

~~R<sup>4a</sup>, at each occurrence, is independently selected from  
H, F, Cl, Br, I, CF<sub>3</sub>,  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
phenyl substituted with 0-3 R<sup>4b</sup>, or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>;~~

~~R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>,  
S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub>  
haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;~~

~~R<sup>5</sup> is H, OR<sup>14</sup>;  
C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>5b</sup>;~~

~~R<sup>5a</sup> is H, methyl, ethyl, propyl, or butyl;~~

~~R<sup>5b</sup>, at each occurrence, is independently selected from:  
H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, Cl, F, Br, I,  
=O;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>5c</sup>;  
phenyl substituted with 0-3 R<sup>5c</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and~~

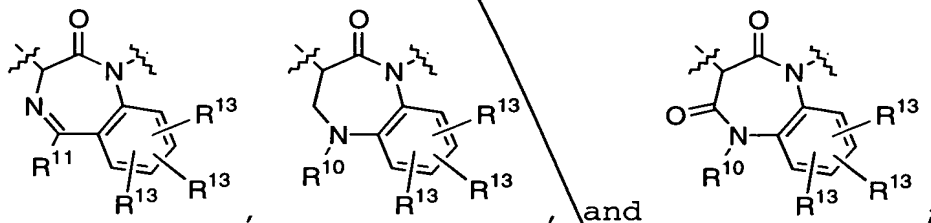
~~sub~~ sulphur, wherein said 5 to 6 membered heterocycle is substituted with 0-3 R<sup>5c</sup>;

~~R<sup>5c</sup>, at each occurrence, is independently selected from H, OH, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;~~

~~R<sup>6</sup> is H;~~

~~R<sup>7</sup>, at each occurrence, is independently selected from H, F, CF<sub>3</sub>, methyl, and ethyl;~~

~~Ring B is selected from~~



~~R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>;~~

~~C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>10a</sup>;~~

~~phenyl substituted with 0-4 R<sup>10b</sup>;~~

~~C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or~~

~~5 to 6 membered heterocycle containing 1 to 4~~

~~heteroatoms selected from nitrogen, oxygen, and~~

~~sulphur, wherein said 5 to 6 membered heterocycle is substituted with 0-3 R<sup>10b</sup>;~~

~~R<sup>10a</sup> is selected from H, C<sub>1</sub>-C<sub>4</sub> alkyl, OR<sup>14</sup>, Cl, F, Br, I, =O, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-4 R<sup>10b</sup>;~~

~~R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>, NR<sup>15</sup>R<sup>16</sup>, or CF<sub>3</sub>;~~



B-5  
Sub  
C7  
R<sup>11</sup> is selected from

H, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, NR<sup>18</sup>R<sup>19</sup>, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>, CF<sub>3</sub>;  
C<sub>1</sub>-C<sub>6</sub> alkyl optionally substituted with 0-3 R<sup>11a</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-3 R<sup>11b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>11b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>11b</sup>;

R<sup>11a</sup>, at each occurrence, is independently selected from H,  
C<sub>1</sub>-C<sub>4</sub> alkyl, OR<sup>14</sup>, F, =O, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl  
substituted with 0-3 R<sup>11b</sup>;

R<sup>11b</sup>, at each occurrence, is independently selected from H,  
OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub>  
haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

Z is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-4 R<sup>12b</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>12b</sup>;

R<sup>12</sup>, at each occurrence, is independently selected from  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-4 R<sup>12b</sup>; or  
5 to 10 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 10 membered heterocycle is  
substituted with 0-3 R<sup>12b</sup>;

B7C  
R<sup>12b</sup>, at each occurrence, is independently selected from  
H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>,  
S(=O)<sub>2</sub>CH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>3</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and  
C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from  
H, OH, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, Cl, F, Br, I, CN, NO<sub>2</sub>,  
NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

Sub  
C7  
R<sup>14</sup> is H, phenyl, benzyl, C<sub>1</sub>-C<sub>4</sub> alkyl, or C<sub>2</sub>-C<sub>4</sub> alkoxyalkyl;

R<sup>15</sup>, at each occurrence, is independently selected from H, C<sub>1</sub>-  
C<sub>4</sub> alkyl, benzyl, phenethyl, (C<sub>1</sub>-C<sub>4</sub> alkyl)-C(=O)-, and  
(C<sub>1</sub>-C<sub>4</sub> alkyl)-S(=O)<sub>2</sub>-;

R<sup>16</sup>, at each occurrence, is independently selected from  
H, OH, C<sub>1</sub>-C<sub>4</sub> alkyl, benzyl, phenethyl,  
(C<sub>1</sub>-C<sub>4</sub> alkyl)-C(=O)-, and (C<sub>1</sub>-C<sub>4</sub> alkyl)-S(=O)<sub>2</sub>-;

R<sup>17</sup> is H, methyl, ethyl, propyl, butyl, methoxymethyl,  
ethoxymethyl, methoxyethyl, ethoxyethyl,  
phenyl substituted by 0-3 R<sup>17a</sup>, or  
-CH<sub>2</sub>-phenyl substituted by 0-3 R<sup>17a</sup>;

R<sup>17a</sup> is H, methyl, methoxy, -OH, F, Cl, CF<sub>3</sub>, or OCF<sub>3</sub>;

R<sup>18</sup>, at each occurrence, is independently selected from  
H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and  
phenethyl; and

R<sup>19</sup>, at each occurrence, is independently selected from  
H, methyl, and ethyl;

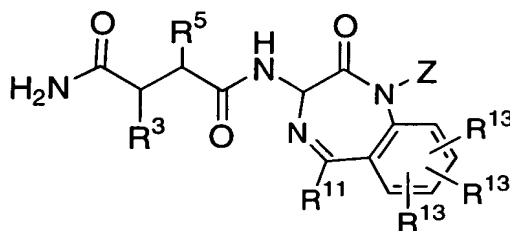
provided, when R<sup>13</sup> is H,

then Z is butyl substituted with 1-3 R<sup>12</sup>;

C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 1-3 R<sup>12</sup>; or

C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 1-3 R<sup>12</sup>.

16. (Twice Amended) A compound according to Claim 14 of Formula (Ic):



(Ic)

or a pharmaceutically acceptable salt thereof  
wherein

R<sup>3</sup> is R<sup>4</sup>,

R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>4a</sup>, or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>;

R<sup>4a</sup> is selected from

H, F, CF<sub>3</sub>,

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,

phenyl substituted with 0-3 R<sup>4b</sup>, or

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

sulphur, wherein said 5 to 6 membered heterocycle is

substituted with 0-3 R<sup>4b</sup>; wherein said 5 to 6 membered

heterocycle is selected from pyridinyl, pyrimidinyl,

triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,

piperazinyl, piperidinyl, pyrazolyl, imidazolyl,

oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy,  
C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

*B#5*  
*Sub*  
*C8*  
~~R<sup>5</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>5b</sup>;~~

~~R<sup>5b</sup> is selected from:~~

~~H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, =O;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-2 R<sup>5c</sup>;  
phenyl substituted with 0-3 R<sup>5c</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>; wherein said 5 to 6 membered  
heterocycle is selected from pyridinyl, pyrimidinyl,  
triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,  
piperazinyl, piperidinyl, pyrazolyl, imidazolyl,  
oxazolyl, isoxazolyl, and tetrazolyl;~~

~~R<sup>5c</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy,  
C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;~~

~~R<sup>11</sup> is selected from~~

~~H, NR<sup>18</sup>R<sup>19</sup>, CF<sub>3</sub>;  
C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>11a</sup>;  
phenyl substituted with 0-3 R<sup>11b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>11b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>11b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;~~

~~R<sup>11a</sup> is selected from H, C<sub>1</sub>-C<sub>4</sub> alkyl, OR<sup>14</sup>, F, =O, NR<sup>15</sup>R<sup>16</sup>,  
CF<sub>3</sub>, or phenyl substituted with 0-3 R<sup>11b</sup>;~~

~~R<sup>11b</sup>, at each occurrence, is independently selected from H,  
OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, methyl, ethyl, propyl, butyl,  
methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub>  
haloalkoxy;~~

~~Z is C<sub>1</sub>-C<sub>3</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>3</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>3</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>12b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>12b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;~~

~~R<sup>12</sup>, at each occurrence, is independently selected from  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>12b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>12b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;~~

~~R<sup>12b</sup>, at each occurrence, is independently selected from~~

H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

~~B5~~  
Sub C8  
R<sup>13</sup>, at each occurrence, is independently selected from H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, Cl, F, Br, CN, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

R<sup>15</sup>, at each occurrence, is independently selected from H, methyl, ethyl, propyl, and butyl;

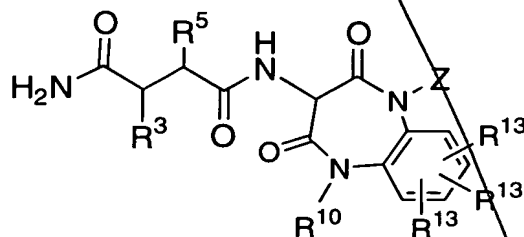
R<sup>16</sup>, at each occurrence, is independently selected from H, OH, methyl, ethyl, propyl, butyl, benzyl, phenethyl, methyl-C(=O)-, ethyl-C(=O)-, methyl-S(=O)<sub>2</sub>-, and ethyl-S(=O)<sub>2</sub>-;

R<sup>18</sup>, at each occurrence, is independently selected from H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and phenethyl; and

R<sup>19</sup>, at each occurrence, is independently selected from H, methyl, and ethyl;

provided, when R<sup>13</sup> is H,  
then Z is C<sub>2</sub>-C<sub>3</sub> alkenyl substituted with 1-3 R<sup>12</sup>; or  
C<sub>2</sub>-C<sub>3</sub> alkynyl substituted with 1-3 R<sup>12</sup>.

~~B5~~  
Sub C9  
18. (Twice Amended) A compound according to Claim 14 of Formula (Ie):



(Ie)

or a pharmaceutically acceptable salt thereof wherein:

*36*  
*Sub*  
*C9*  
R<sup>3</sup> is R<sup>4</sup>,

R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>4a</sup>, or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>;

R<sup>4a</sup> is selected from

H, F, CF<sub>3</sub>,

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,

phenyl substituted with 0-3 R<sup>4b</sup>, or

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

sulphur, wherein said 5 to 6 membered heterocycle is

substituted with 0-3 R<sup>4b</sup>; wherein said 5 to 6 membered

heterocycle is selected from pyridinyl, pyrimidinyl,

triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,

piperazinyl, piperidinyl, pyrazolyl, imidazolyl,

oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy,  
C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>5</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>5b</sup>;

R<sup>5b</sup> is selected from:

H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, =O;

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-2 R<sup>5c</sup>;

phenyl substituted with 0-3 R<sup>5c</sup>; or

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and

85  
sulphur, wherein said 5 to 6 membered heterocycle is substituted with 0-3 R<sup>5c</sup>; wherein said 5 to 6 membered heterocycle is selected from pyridinyl, pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl, piperazinyl, piperidinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>5c</sup>, at each occurrence, is independently selected from H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>;  
C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>10a</sup>;  
phenyl substituted with 0-4 R<sup>10b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 6 membered heterocycle is substituted with 0-3 R<sup>10b</sup>; wherein said 5 to 6 membered heterocycle is selected from pyridinyl, pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl, piperazinyl, piperidinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>10a</sup> is selected from H, methyl, ethyl, propyl, butyl, OR<sup>14</sup>, Cl, F, =O, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-4 R<sup>10b</sup>;

R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, Cl, F, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

Z is C<sub>1</sub>-C<sub>3</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>3</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>3</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;



36  
Sub  
C9

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>12b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>12b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>12</sup>, at each occurrence, is independently selected from  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>12b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>12b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>12b</sup>, at each occurrence, is independently selected from  
H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>,  
S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from  
H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
Cl, F, Br, CN, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

R<sup>15</sup>, at each occurrence, is independently selected from H,  
methyl, ethyl, propyl, and butyl;

R<sup>16</sup>, at each occurrence, is independently selected from

H, OH, methyl, ethyl, propyl, butyl, benzyl, phenethyl,  
methyl-C(=O)-, ethyl-C(=O)-,  
methyl-S(=O)<sub>2</sub>-, and ethyl-S(=O)<sub>2</sub>-;

30  
Sub  
C9  
R<sup>17</sup> is H, methyl, ethyl, propyl, butyl, methoxymethyl,  
ethoxymethyl, methoxyethyl, ethoxyethyl,  
phenyl substituted by 0-3 R<sup>17a</sup>, or  
-CH<sub>2</sub>-phenyl substituted by 0-3 R<sup>17a</sup>;

R<sup>17a</sup> is H, methyl, methoxy, -OH, F, Cl, CF<sub>3</sub>, or OCF<sub>3</sub>;

R<sup>18</sup>, at each occurrence, is independently selected from  
H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and  
phenethyl; and

R<sup>19</sup>, at each occurrence, is independently selected from  
H, methyl, and ethyl;

provided, when R<sup>13</sup> is H,  
then Z is C<sub>2</sub>-C<sub>3</sub> alkenyl substituted with 1-3 R<sup>12</sup>; or  
C<sub>2</sub>-C<sub>3</sub> alkynyl substituted with 1-3 R<sup>12</sup>.

20. (Twice Amended) A compound according to one of Claims 16,  
18, or 26 wherein:

30  
Sub  
C10  
R<sup>3</sup> is -CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>,  
-CH(CH<sub>3</sub>)<sub>2</sub>, -CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH(CH<sub>3</sub>)<sub>2</sub>,  
-CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>,  
-CH=CH<sub>2</sub>, -CH<sub>2</sub>CH=CH<sub>2</sub>, -CH<sub>2</sub>C(CH<sub>3</sub>)=CH<sub>2</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH=CH<sub>2</sub>,  
cis-CH<sub>2</sub>CH=CH(CH<sub>3</sub>),  
trans-CH<sub>2</sub>CH=CH(CH<sub>3</sub>),  
-C≡CH, -CH<sub>2</sub>C≡CH, -CH<sub>2</sub>C≡C(CH<sub>3</sub>),  
cyclopropyl-CH<sub>2</sub>-, cyclobutyl-CH<sub>2</sub>-, cyclopentyl-CH<sub>2</sub>-,  
cyclohexyl-CH<sub>2</sub>-, cyclopropyl-CH<sub>2</sub>CH<sub>2</sub>-,  
cyclobutyl-CH<sub>2</sub>CH<sub>2</sub>-, cyclopentyl-CH<sub>2</sub>CH<sub>2</sub>-,  
cyclohexyl-CH<sub>2</sub>CH<sub>2</sub>-, phenyl-CH<sub>2</sub>-,

64  
5.10  
C10

(2-F-phenyl)CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>-, (4-F-phenyl)CH<sub>2</sub>-,  
(2-Cl-phenyl)CH<sub>2</sub>-, (3-Cl-phenyl)CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>-,  
(2,3-diF-phenyl)CH<sub>2</sub>-, (2,4-diF-phenyl)CH<sub>2</sub>-,  
(2,5-diF-phenyl)CH<sub>2</sub>-, (2,6-diF-phenyl)CH<sub>2</sub>-,  
(3,4-diF-phenyl)CH<sub>2</sub>-, (3,5-diF-phenyl)CH<sub>2</sub>-,  
(2,3-diCl-phenyl)CH<sub>2</sub>-, (2,4-diCl-phenyl)CH<sub>2</sub>-,  
(2,5-diCl-phenyl)CH<sub>2</sub>-, (2,6-diCl-phenyl)CH<sub>2</sub>-,  
(3,4-diCl-phenyl)CH<sub>2</sub>-, (3,5-diCl-phenyl)CH<sub>2</sub>-,  
(3-F-4-Cl-phenyl)CH<sub>2</sub>-, (3-F-5-Cl-phenyl)CH<sub>2</sub>-,  
(3-Cl-4-F-phenyl)CH<sub>2</sub>-, phenyl-CH<sub>2</sub>CH<sub>2</sub>-,  
(2-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,3-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,4-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,5-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,6-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3,4-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3,5-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,3-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,4-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,5-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,6-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3,4-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3,5-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-F-4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, or (3-F-5-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,

R<sup>5</sup> is -CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH(CH<sub>3</sub>)<sub>2</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>,  
-CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH(CH<sub>3</sub>)<sub>2</sub>, -CH<sub>2</sub>C(CH<sub>3</sub>)<sub>3</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CH<sub>2</sub>CH(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>3</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH(CH<sub>3</sub>)<sub>2</sub>, -CH(CH<sub>2</sub>CH<sub>3</sub>)<sub>2</sub>, -CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>,  
-CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CF<sub>3</sub>, -CH=CH<sub>2</sub>, -CH<sub>2</sub>CH=CH<sub>2</sub>,  
-CH=CHCH<sub>3</sub>, cis-CH<sub>2</sub>CH=CH(CH<sub>3</sub>), trans-CH<sub>2</sub>CH=CH(CH<sub>3</sub>),  
trans-CH<sub>2</sub>CH=CH(C<sub>6</sub>H<sub>5</sub>), -CH<sub>2</sub>CH=C(CH<sub>3</sub>)<sub>2</sub>, cis-CH<sub>2</sub>CH=CHCH<sub>2</sub>CH<sub>3</sub>,  
trans-CH<sub>2</sub>CH=CHCH<sub>2</sub>CH<sub>3</sub>, cis-CH<sub>2</sub>CH<sub>2</sub>CH=CH(CH<sub>3</sub>),  
trans-CH<sub>2</sub>CH<sub>2</sub>CH=CH(CH<sub>3</sub>), trans-CH<sub>2</sub>CH=CHCH<sub>2</sub>(C<sub>6</sub>H<sub>5</sub>),  
-C≡CH, -CH<sub>2</sub>C≡CH, -CH<sub>2</sub>C≡C(CH<sub>3</sub>), -CH<sub>2</sub>C≡C(C<sub>6</sub>H<sub>5</sub>),  
-CH<sub>2</sub>CH<sub>2</sub>C≡CH, -CH<sub>2</sub>CH<sub>2</sub>C≡C(CH<sub>3</sub>), -CH<sub>2</sub>CH<sub>2</sub>C≡C(C<sub>6</sub>H<sub>5</sub>),  
cyclopropyl-CH<sub>2</sub>-, cyclobutyl-CH<sub>2</sub>-, cyclopentyl-CH<sub>2</sub>-,  
cyclohexyl-CH<sub>2</sub>-, (2-CH<sub>3</sub>-cyclopropyl)CH<sub>2</sub>-,  
(3-CH<sub>3</sub>-cyclobutyl)CH<sub>2</sub>-,  
cyclopropyl-CH<sub>2</sub>CH<sub>2</sub>-, cyclobutyl-CH<sub>2</sub>CH<sub>2</sub>-,  
cyclopentyl-CH<sub>2</sub>CH<sub>2</sub>-, cyclohexyl-CH<sub>2</sub>CH<sub>2</sub>-,

*B6*  
*Sub*  
*C10*  
(2-CH<sub>3</sub>-cyclopropyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-CH<sub>3</sub>-cyclobutyl)CH<sub>2</sub>CH<sub>2</sub>-,  
phenyl-CH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>-,  
(4-F-phenyl)CH<sub>2</sub>-, furanyl-CH<sub>2</sub>-, thienyl-CH<sub>2</sub>-,  
pyridyl-CH<sub>2</sub>-, 1-imidazolyl-CH<sub>2</sub>-, oxazolyl-CH<sub>2</sub>-,  
isoxazolyl-CH<sub>2</sub>-,  
phenyl-CH<sub>2</sub>CH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, furanyl-CH<sub>2</sub>CH<sub>2</sub>-, thienyl-CH<sub>2</sub>CH<sub>2</sub>-,  
pyridyl-CH<sub>2</sub>CH<sub>2</sub>-, 1-imidazolyl-CH<sub>2</sub>CH<sub>2</sub>-, oxazolyl-CH<sub>2</sub>CH<sub>2</sub>-,  
isoxazolyl-CH<sub>2</sub>CH<sub>2</sub>-;

Z is phenyl, 2-F-phenyl, 3-F-phenyl, 4-F-phenyl,  
2-Cl-phenyl, 3-Cl-phenyl, 4-Cl-phenyl, 2,3-diF-phenyl,  
2,4-diF-phenyl, 2,5-diF-phenyl, 2,6-diF-phenyl,  
3,4-diF-phenyl, 3,5-diF-phenyl, 2,3-diCl-phenyl,  
2,4-diCl-phenyl, 2,5-diCl-phenyl, 2,6-diCl-phenyl,  
3,4-diCl-phenyl, 3,5-diCl-phenyl, 3-F-4-Cl-phenyl,  
3-F-5-Cl-phenyl, 3-Cl-4-F-phenyl, 2-MeO-phenyl,  
3-MeO-phenyl, 4-MeO-phenyl, 2-Me-phenyl, 3-Me-phenyl,  
4-Me-phenyl, 2-MeS-phenyl, 3-MeS-phenyl, 4-MeS-phenyl,  
2-CF<sub>3</sub>O-phenyl, 3-CF<sub>3</sub>O-phenyl, 4-CF<sub>3</sub>O-phenyl,  
furanyl, thienyl, pyridyl, 2-Me-pyridyl, 3-Me-pyridyl,  
4-Me-pyridyl, 1-imidazolyl, oxazolyl, isoxazolyl,  
cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl,  
N-piperidinyl,  
phenyl-CH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>-,  
(4-F-phenyl)CH<sub>2</sub>-, (2-Cl-phenyl)CH<sub>2</sub>-, (3-Cl-phenyl)CH<sub>2</sub>-, (4-  
Cl-phenyl)CH<sub>2</sub>-, (2,3-diF-phenyl)CH<sub>2</sub>-,  
(2,4-diF-phenyl)CH<sub>2</sub>-, (2,5-diF-phenyl)CH<sub>2</sub>-,  
(2,6-diF-phenyl)CH<sub>2</sub>-, (3,4-diF-phenyl)CH<sub>2</sub>-,  
(3,5-diF-phenyl)CH<sub>2</sub>-, (2,3-diCl-phenyl)CH<sub>2</sub>-,  
(2,4-diCl-phenyl)CH<sub>2</sub>-, (2,5-diCl-phenyl)CH<sub>2</sub>-,  
(2,6-diCl-phenyl)CH<sub>2</sub>-, (3,4-diCl-phenyl)CH<sub>2</sub>-,  
(3,5-diCl-phenyl)CH<sub>2</sub>-, (3-F-4-Cl-phenyl)CH<sub>2</sub>-,  
(3-F-5-Cl-phenyl)CH<sub>2</sub>-, (3-Cl-4-F-phenyl)CH<sub>2</sub>-,  
(2-MeO-phenyl)CH<sub>2</sub>-, (3-MeO-phenyl)CH<sub>2</sub>-,  
(4-MeO-phenyl)CH<sub>2</sub>-, (2-Me-phenyl)CH<sub>2</sub>-,  
(3-Me-phenyl)CH<sub>2</sub>-, (4-Me-phenyl)CH<sub>2</sub>-.

BK  
Sub  
C10  
(2-MeS-phenyl)CH<sub>2</sub>-, (3-MeS-phenyl)CH<sub>2</sub>-,  
4-MeS-phenyl)CH<sub>2</sub>-, (2-CF<sub>3</sub>O-phenyl)CH<sub>2</sub>-,  
(3-CF<sub>3</sub>O-phenyl)CH<sub>2</sub>-, (4-CF<sub>3</sub>O-phenyl)CH<sub>2</sub>-,  
(furanyl)CH<sub>2</sub>-, (thienyl)CH<sub>2</sub>-, (pyridyl)CH<sub>2</sub>-,  
(2-Me-pyridyl)CH<sub>2</sub>-, (3-Me-pyridyl)CH<sub>2</sub>-,  
(4-Me-pyridyl)CH<sub>2</sub>-, (1-imidazolyl)CH<sub>2</sub>-,  
(oxazolyl)CH<sub>2</sub>-, (isoxazolyl)CH<sub>2</sub>-,  
(cyclopropyl)CH<sub>2</sub>-, (cyclobutyl)CH<sub>2</sub>-, (cyclopentyl)CH<sub>2</sub>-,  
(cyclohexyl)CH<sub>2</sub>-, (N-piperidiny)CH<sub>2</sub>-,

phenyl-CH<sub>2</sub>CH<sub>2</sub>-, (phenyl)<sub>2</sub>CHCH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,3-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,4-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,5-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,6-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3,4-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3,5-diF-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,3-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,4-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2,5-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2,6-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3,4-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3,5-diCl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-F-4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-F-5-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-Cl-4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2-MeO-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-MeO-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-MeO-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2-Me-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-Me-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (4-Me-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2-MeS-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-MeS-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-MeS-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (2-CF<sub>3</sub>O-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(3-CF<sub>3</sub>O-phenyl)CH<sub>2</sub>CH<sub>2</sub>-, (4-CF<sub>3</sub>O-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(furanyl)CH<sub>2</sub>CH<sub>2</sub>-, (thienyl)CH<sub>2</sub>CH<sub>2</sub>-, (pyridyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(2-Me-pyridyl)CH<sub>2</sub>CH<sub>2</sub>-, (3-Me-pyridyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(4-Me-pyridyl)CH<sub>2</sub>CH<sub>2</sub>-, (imidazolyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(oxazolyl)CH<sub>2</sub>CH<sub>2</sub>-, (isoxazolyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(cyclopropyl)CH<sub>2</sub>CH<sub>2</sub>-, (cyclobutyl)CH<sub>2</sub>CH<sub>2</sub>-,  
(cyclopentyl)CH<sub>2</sub>CH<sub>2</sub>-, (cyclohexyl)CH<sub>2</sub>CH<sub>2</sub>-, or  
(N-piperidiny)CH<sub>2</sub>CH<sub>2</sub>-;

R<sup>10</sup> is H, methyl, ethyl, phenyl, benzyl, phenethyl,  
4-F-phenyl, (4-F-phenyl)CH<sub>2</sub>-, (4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,

4-Cl-phenyl, (4-Cl-phenyl)CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CH<sub>3</sub>-phenyl, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>-, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CF<sub>3</sub>-phenyl, (4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>-, or  
(4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-;

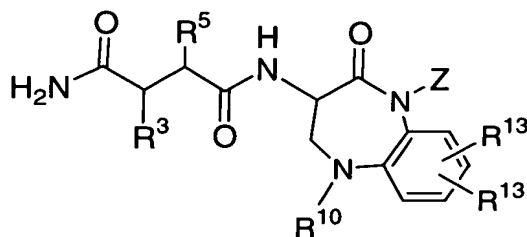
B8  
Sub  
C10  
R<sup>11</sup>, at each occurrence, is independently selected from  
H, =O, methyl, ethyl, phenyl, benzyl, phenethyl,  
4-F-phenyl, (4-F-phenyl)CH<sub>2</sub>-, (4-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
3-F-phenyl, (3-F-phenyl)CH<sub>2</sub>-, (3-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
2-F-phenyl, (2-F-phenyl)CH<sub>2</sub>-, (2-F-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-Cl-phenyl, (4-Cl-phenyl)CH<sub>2</sub>-, (4-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
3-Cl-phenyl, (3-Cl-phenyl)CH<sub>2</sub>-, (3-Cl-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CH<sub>3</sub>-phenyl, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>-, (4-CH<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
3-CH<sub>3</sub>-phenyl, (3-CH<sub>3</sub>-phenyl)CH<sub>2</sub>-, (3-CH<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
4-CF<sub>3</sub>-phenyl, (4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>-, (4-CF<sub>3</sub>-phenyl)CH<sub>2</sub>CH<sub>2</sub>-,  
pyrid-2-yl, pyrid-3-yl, or pyrid-4-yl, and

R<sup>13</sup>, at each occurrence, is independently selected from  
H, F, Cl, OH, -CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>3</sub>, -OCH<sub>3</sub>, or -CF<sub>3</sub>.

B8  
Sub  
C11  
22. A pharmaceutical composition comprising a compound of  
Claim 1 and a pharmaceutically acceptable carrier.

23. (Amended) A method for the treatment of Alzheimer's  
Disease production comprising administering to a host in need  
of such treatment a therapeutically effective amount of a  
compound of Claim 1.

B8  
25. (New) A compound according to Claim 4 of Formula (Ig):



(Ig)

or a pharmaceutically acceptable salt thereof wherein:

*B 8*  
R<sup>3</sup> is R<sup>4</sup>,

R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>4a</sup>, or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>;

R<sup>4a</sup>, at each occurrence, is independently selected from  
H, F, CF<sub>3</sub>,  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,  
phenyl substituted with 0-3 R<sup>4b</sup>, or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>; wherein said 5 to 6 membered  
heterocycle is selected from pyridinyl, pyrimidinyl,  
triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,  
piperazinyl, piperidinyl, pyrazolyl, imidazolyl,  
oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy,  
C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>5</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>5b</sup>;

R<sup>5b</sup> is selected from:

H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, =O;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-2 R<sup>5c</sup>;  
phenyl substituted with 0-3 R<sup>5c</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is

B9 substituted with 0-3 R<sup>5c</sup>; wherein said 5 to 6 membered heterocycle is selected from pyridinyl, pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl, piperazinyl, piperidinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>5c</sup>, at each occurrence, is independently selected from H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>;  
C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>10a</sup>;  
phenyl substituted with 0-4 R<sup>10b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 6 membered heterocycle is substituted with 0-3 R<sup>10b</sup>; wherein said 5 to 6 membered heterocycle is selected from pyridinyl, pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl, piperazinyl, piperidinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>10a</sup> is selected from H, methyl, ethyl, propyl, butyl, OR<sup>14</sup>, Cl, F, =O, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-4 R<sup>10b</sup>;


R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, Cl, F, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

Z is H;

C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-3 R<sup>12a</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-3 R<sup>12a</sup>; or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-3 R<sup>12a</sup>;



R<sup>12a</sup>, at each occurrence, is independently selected from  
H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>,  
S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

 R<sup>13</sup>, at each occurrence, is independently selected from  
H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
Cl, F, Br, CN, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

R<sup>15</sup>, at each occurrence, is independently selected from H,  
methyl, ethyl, propyl, and butyl;

R<sup>16</sup>, at each occurrence, is independently selected from  
H, OH, methyl, ethyl, propyl, butyl, benzyl, phenethyl,  
methyl-C(=O)-, ethyl-C(=O)-,  
methyl-S(=O)<sub>2</sub>-, and ethyl-S(=O)<sub>2</sub>-;

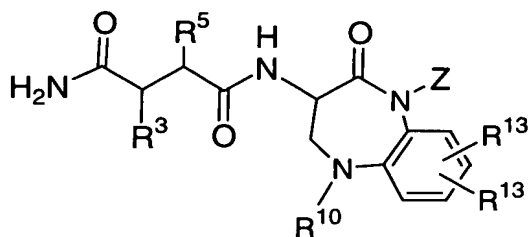
R<sup>17</sup> is H, methyl, ethyl, propyl, butyl, methoxymethyl,  
ethoxymethyl, methoxyethyl, ethoxyethyl,  
phenyl substituted by 0-3 R<sup>17a</sup>, or  
-CH<sub>2</sub>-phenyl substituted by 0-3 R<sup>17a</sup>;

R<sup>17a</sup> is H, methyl, methoxy, -OH, F, Cl, CF<sub>3</sub>, or OCF<sub>3</sub>;

R<sup>18</sup>, at each occurrence, is independently selected from  
H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and  
phenethyl; and

R<sup>19</sup>, at each occurrence, is independently selected from  
H, methyl, and ethyl.

**26.** (New) A compound according to Claim 14 of Formula (Ig):



(Ig)

or a pharmaceutically acceptable salt thereof wherein:

R<sup>3</sup> is R<sup>4</sup>,

R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>4a</sup>,  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>4a</sup>, or  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>4a</sup>;

R<sup>4a</sup> is selected from

H, F, CF<sub>3</sub>,

C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>4b</sup>,

phenyl substituted with 0-3 R<sup>4b</sup>, or

5 to 6 membered heterocycle containing 1 to 4

heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>4b</sup>; wherein said 5 to 6 membered  
heterocycle is selected from pyridinyl, pyrimidinyl,  
triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,  
piperazinyl, piperidinyl, pyrazolyl, imidazolyl,  
oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>4b</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy,  
C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>5</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkenyl substituted with 0-1 R<sup>5b</sup>;  
C<sub>2</sub>-C<sub>4</sub> alkynyl substituted with 0-1 R<sup>5b</sup>;

B<sup>8</sup>  
R<sup>5b</sup> is selected from:


H, methyl, ethyl, propyl, butyl, CF<sub>3</sub>, OR<sup>14</sup>, =O;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-2 R<sup>5c</sup>;  
phenyl substituted with 0-3 R<sup>5c</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>5c</sup>; wherein said 5 to 6 membered  
heterocycle is selected from pyridinyl, pyrimidinyl,  
triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl,  
piperazinyl, piperidinyl, pyrazolyl, imidazolyl,  
oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>5c</sup>, at each occurrence, is independently selected from H, OH,  
Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>,  
methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy,  
C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>10</sup> is H, C(=O)R<sup>17</sup>, C(=O)OR<sup>17</sup>;  
C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with 0-1 R<sup>10a</sup>;  
phenyl substituted with 0-4 R<sup>10b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>10b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4  
heteroatoms selected from nitrogen, oxygen, and  
sulphur, wherein said 5 to 6 membered heterocycle is  
substituted with 0-3 R<sup>10b</sup>; wherein said 5 to 6  
membered heterocycle is selected from pyridinyl,  
pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl,  
pyrrolyl, piperazinyl, piperidinyl, pyrazolyl,  
imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>10a</sup> is selected from H, methyl, ethyl, propyl, butyl, OR<sup>14</sup>,  
Cl, F, =O, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, or phenyl substituted with 0-4  
R<sup>10b</sup>;

R<sup>10b</sup>, at each occurrence, is independently selected from H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, Cl, F, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

 Z is C<sub>1</sub>-C<sub>3</sub> alkyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>3</sub> alkenyl substituted with 1-3 R<sup>12</sup>;  
C<sub>2</sub>-C<sub>3</sub> alkynyl substituted with 1-3 R<sup>12</sup>;  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>12b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 6 membered heterocycle is substituted with 0-3 R<sup>12b</sup>; wherein said 5 to 6 membered heterocycle is selected from pyridinyl, pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl, piperazinyl, piperidinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>12</sup>, at each occurrence, is independently selected from  
C<sub>6</sub>-C<sub>10</sub> aryl substituted with 0-4 R<sup>12b</sup>;  
C<sub>3</sub>-C<sub>6</sub> carbocycle substituted with 0-3 R<sup>12b</sup>; or  
5 to 6 membered heterocycle containing 1 to 4 heteroatoms selected from nitrogen, oxygen, and sulphur, wherein said 5 to 6 membered heterocycle is substituted with 0-3 R<sup>12b</sup>; wherein said 5 to 6 membered heterocycle is selected from pyridinyl, pyrimidinyl, triazinyl, furanyl, thienyl, thiazolyl, pyrrolyl, piperazinyl, piperidinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, and tetrazolyl;

R<sup>12b</sup>, at each occurrence, is independently selected from  
H, OH, Cl, F, NR<sup>15</sup>R<sup>16</sup>, CF<sub>3</sub>, acetyl, SCH<sub>3</sub>, S(=O)CH<sub>3</sub>, S(=O)<sub>2</sub>CH<sub>3</sub>, methyl, ethyl, propyl, butyl, methoxy, ethoxy, propoxy, C<sub>1</sub>-C<sub>2</sub> haloalkyl, and C<sub>1</sub>-C<sub>2</sub> haloalkoxy;

R<sup>13</sup>, at each occurrence, is independently selected from

H, OH, methyl, ethyl, propyl, butyl, methoxy, ethoxy,  
Cl, F, Br, CN, NR<sup>15</sup>R<sup>16</sup>, and CF<sub>3</sub>;

R<sup>14</sup> is H, phenyl, benzyl, methyl, ethyl, propyl, or butyl;

*BA* R<sup>15</sup>, at each occurrence, is independently selected from H,  
methyl, ethyl, propyl, and butyl;

R<sup>16</sup>, at each occurrence, is independently selected from  
H, OH, methyl, ethyl, propyl, butyl, benzyl, phenethyl,  
methyl-C(=O)-, ethyl-C(=O)-,  
methyl-S(=O)<sub>2</sub>-, and ethyl-S(=O)<sub>2</sub>-;

R<sup>17</sup> is H, methyl, ethyl, propyl, butyl, methoxymethyl,  
ethoxymethyl, methoxyethyl, ethoxyethyl,  
phenyl substituted by 0-3 R<sup>17a</sup>, or  
-CH<sub>2</sub>-phenyl substituted by 0-3 R<sup>17a</sup>;

R<sup>17a</sup> is H, methyl, methoxy, -OH, F, Cl, CF<sub>3</sub>, or OCF<sub>3</sub>;

R<sup>18</sup>, at each occurrence, is independently selected from  
H, methyl, ethyl, propyl, butyl, phenyl, benzyl, and  
phenethyl; and

R<sup>19</sup>, at each occurrence, is independently selected from  
H, methyl, and ethyl;

provided, when R<sup>13</sup> is H,  
then Z is C<sub>2</sub>-C<sub>3</sub> alkenyl substituted with 1-3 R<sup>12</sup>; or  
C<sub>2</sub>-C<sub>3</sub> alkynyl substituted with 1-3 R<sup>12</sup>. --

---